K063066

AUG - 3 2007

510(k) SUMMARY

A. Manufacturer:	National Display Systems, LLC 16245 Vineyard Boulevard Morgan Hill, CA 95037 USA
B. Submitted By:	Ron Hansen Chief Technical Officer National Display Systems, LLC
C. Date of Preparation:	August 24, 2006
D. Contact Information:	Tel: 408-776-0085 Fax: 408-776-9878
E. Classification:	System, image processing, radiological
F. Common Name:	Monitor, display, and others
G. Proprietary Name:	AXIS-V 5MP Monochrome Display
H. Classification Number:	21 CFR 892.2050/Procode 90LLZ
I. Substantial Equivalence:	Coronis 5MP (Barco) K042221 RadiForce G51 (Eizo Nanao) K042755
J. Device Description:	The AXIS-V 5MP Monochrome Display is a diagnostic display.
K. Intended Use:	The AXIS-V 5MP Radiological Monochrome Medical Display is intended to be used to display and view digital images, including digital mammography, for review and analysis by trained medical practitioners.
L. Technological Characteristics:	The AXIS-V 5MP Monochrome Display is a high-resolution Liquid Crystal Display (LCD) with electronic capabilities used for the review and analysis of high-resolution medical images by trained medical practitioners.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
9200 Corporate Blvd.
Rockville MD 20850

AUG - 3 2007

National Display Systems
% Ms. Denise Leung Klinker
Staff Engineer / 510(k) Reviewer
Underwriters Laboratories, Inc.
455 East Trimble Road
SAN JOSE CA 95131

Re: K063066
Trade/Device Name: AXIS-V 5 MP Radiology Medical Display
Regulation Number: 21 CFR 892.2050
Regulation Name: Picture archiving and communications system
Regulatory Class: II
Product Code: LLZ
Dated: July 23, 2007
Received: July 24, 2007

Dear Ms. Klinker:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Office of Compliance at one of the following numbers, based on the regulation number at the top of this letter:

21 CFR 876.xxx	(Gastroenterology/Renal/Urology	240-276-0115
21 CFR 884.xxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 894.xxx	(Radiology)	240-276-0120
Other		240-276-0100

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (240) 276-3150
or at its Internet address http://www.fda.gov/cdrh/industry/support/index.html.

Sincerely yours,

Nancy C Brogdon

Nancy C. Brogdon
Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): ~~N/A~~ K063066

Device Name: AXIS-V 5MP Radiology Medical Display

Indications for Use:

The AXIS-V 5MP Radiological Monochrome Medical Display is intended to be used to display and view digital images, including digital mammography, for review and analysis by trained medical practitioners.

Prescription Use __X__ AND/OR Over-the-Counter Use _____
(Part 21 CFR 801 Subpart D) (21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIE LINE – CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

(Division Sign-Off)
Division of Reproductive, Abdominal and
Radiological Devices
510(k) Number K063066